



DCN gene

decorin

Normal Function

The *DCN* gene provides instructions for making a protein called decorin. This protein is a component of the extracellular matrix, which is the intricate lattice of proteins and other molecules that forms in the spaces between cells. Decorin is found in the extracellular matrix of a variety of connective tissues, including skin, tendon, bone, and cartilage. Connective tissues support the body's joints and organs.

Decorin is involved in the organization of proteins called collagens. Collagens strengthen and support connective tissues throughout the body. Collagens also play an important role in the cornea, which is the clear outer covering of the eye. Bundles of collagen called fibrils must be strictly organized for the cornea to be transparent. Decorin ensures that these collagen fibrils are uniformly sized and regularly spaced.

Researchers have proposed several additional functions for decorin. This protein likely helps regulate cell growth and division, the attachment of cells to one another (cell adhesion), and the self-destruction of cells (apoptosis). Studies suggest that decorin plays a role in the formation of new blood vessels (angiogenesis), wound healing, bone development, inflammation, and preventing the growth of cancerous tumors. Decorin also regulates the activity of several growth factors, including transforming growth factor-beta (TGF β). These growth factors control a diverse range of processes important for cell growth.

Health Conditions Related to Genetic Changes

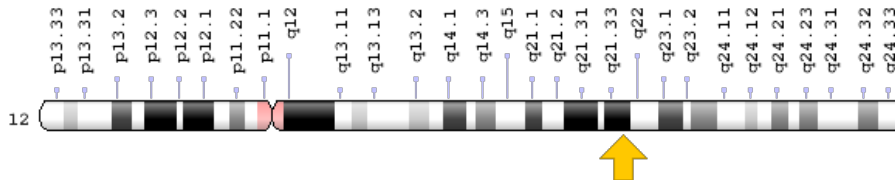
congenital stromal corneal dystrophy

Several mutations in the *DCN* gene have been identified in families with congenital stromal corneal dystrophy. Each of the known mutations leads to the production of an abnormally short version of the decorin protein. This abnormal protein interferes with the organization of collagen fibrils in the cornea. As poorly arranged collagen fibrils accumulate, the cornea becomes cloudy. These corneal changes lead to a loss of sharp vision (reduced visual acuity) and other eye abnormalities related to visual impairment.

Chromosomal Location

Cytogenetic Location: 12q21.33, which is the long (q) arm of chromosome 12 at position 21.33

Molecular Location: base pairs 91,143,277 to 91,183,124 on chromosome 12 (Homo sapiens Annotation Release 108, GRCh38.p7) (NCBI)



Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- bone proteoglycan II
- decorin proteoglycan
- dermatan sulphate proteoglycans II
- DSPG2
- PG40
- PGII
- PGS2
- PGS2_HUMAN
- proteoglycan core protein
- SLRR1B
- small leucine-rich protein 1B

Additional Information & Resources

Educational Resources

- Molecular Biology of the Cell (fourth edition, 2002): Collagens Are the Major Proteins of the Extracellular Matrix
<https://www.ncbi.nlm.nih.gov/books/NBK26810/#A3551>

GeneReviews

- Congenital Stromal Corneal Dystrophy
<https://www.ncbi.nlm.nih.gov/books/NBK2690>

Scientific Articles on PubMed

- PubMed
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28DCN%5BTI%5D%29+OR+%28decorin%5BTI%5D%29%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+720+days%22%5Bdp%5D>

OMIM

- DECORIN
<http://omim.org/entry/125255>

Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology
http://atlasgeneticsoncology.org/Genes/GC_DCN.html
- ClinVar
<https://www.ncbi.nlm.nih.gov/clinvar?term=DCN%5Bgene%5D>
- HGNC Gene Family: Small leucine rich repeat proteoglycans
<http://www.genenames.org/cgi-bin/genefamilies/set/573>
- HGNC Gene Symbol Report
http://www.genenames.org/cgi-bin/gene_symbol_report?q=data/hgnc_data.php&hgnc_id=2705
- NCBI Gene
<https://www.ncbi.nlm.nih.gov/gene/1634>
- UniProt
<http://www.uniprot.org/uniprot/P07585>

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Reviewed: August 2009
Published: March 21, 2017

Lister Hill National Center for Biomedical Communications
U.S. National Library of Medicine
National Institutes of Health
Department of Health & Human Services